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1 **Costs and benefits of sub-lethal Drosophila C Virus infection**

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ABSTRACT

Viruses are major evolutionary drivers of insect immune systems. Much of our knowledge of insect immune responses derives from experimental infections using the fruit fly *Drosophila melanogaster*. Most experiments, however, employ lethal pathogen doses through septic injury, frequently overwhelming host physiology. While this approach has revealed several immune mechanisms, it is less informative about the fitness costs hosts may experience during infection in the wild. Using both systemic and oral infection routes we find that even apparently benign, sub-lethal infections with the horizontally transmitted *Drosophila* C Virus (DCV) can cause significant physiological and behavioral morbidity that is relevant for host fitness. We describe DCV-induced effects on fly reproductive output, digestive health, and locomotor activity, and we find that viral morbidity varies according to the concentration of pathogen inoculum, host genetic background and sex. Notably, sub-lethal DCV infection resulted in a significant increase in fly reproduction, but this effect depended on host genotype. We discuss the relevance of sub-lethal morbidity for *Drosophila* ecology and evolution, and more broadly, we remark on the implications of deleterious and beneficial infections for the evolution of insect immunity.

Key-words: Sub-lethal infection; systemic infection; oral infection; fecundity; locomotor activity; fecal excretion; fitness.

INTRODUCTION

Viral infections are pervasive throughout the living world (Suttle, 2005; Rosario & Breitbart, 2011). Viruses of insects have attracted considerable interest (Miller & Ball, eds, 1998), in part due to their potential role in the bio-control of insect pests (Lacey *et al.*, 2015), and also because insects are vectors of many viral pathogens of plants (Whitfield *et al.*, 2015), animals and humans (Conway *et al.*, 2014). The abundance and diversity of insect viruses, combined with the extensive morbidity and mortality they cause, make viral infections potentially powerful determinants of insect population dynamics and evolution (Dwyer *et al.*, 2004; Obbard *et al.*, 2006; Wilfert *et al.*, 2016).

Much of our knowledge of insect immune responses to viral infections has come from work using the fruit fly *Drosophila melanogaster*, where the focus has been on elucidating the genetics underlying antiviral immunity (Dostert *et al.*, 2005; Huszar & Imler, 2008; Kemp & Imler, 2009; Sabin *et al.*, 2010; Magwire *et al.*, 2012). Several RNA viruses have been described and investigated in this context, including Nora virus (Habayeb *et al.*, 2009), Drosophila A virus (DAV) (Ambrose *et al.*, 2009), Flock House Virus (FHV) (Scotti *et al.*, 1983) and Drosophila C Virus (DCV) (Jousset *et al.*, 1977), a horizontally transmitted ssRNA virus in the *Dicistroviridae* family (Huszar & Imler, 2008). Initial investigations of DCV infection found that it replicates in the fly's reproductive and digestive tissues (Lautié-Harivel & Thomas-Orillard, 1990) and that infection results in accelerated larval development but also causes mortality (Thomas-Orillard, 1984; Gomariz-Zilber *et al.*, 1995). More recent work has shown that systemic infection with elevated concentrations of DCV causes pathology within the fly's

food storage organ, the crop, leading to intestinal obstruction, lower metabolic rate and reduced locomotor activity (Arnold *et al.*, 2013; Chtarbanova *et al.*, 2014). There is also considerable genetic variation in fly survival when challenged systemically with DCV, which appears to be controlled by few genes of large effect (Magwire *et al.*, 2012).

While this level of detail concerning the physiological consequences and the underlying genetics of infection is remarkable, it is important to recognize that our knowledge of viral infections comes almost entirely from experimental infections that challenge model systems, such as *Drosophila*, with artificially high viral concentrations during systemic infections. Even in cases where natural routes of infection have been investigated (Gomariz-Zilber *et al.*, 1995; Ferreira *et al.*, 2014; Stevanovic & Johnson, 2015; Vale & Jardine, 2015), these have often been achieved by using much higher doses than flies are likely to encounter in the wild in order to cause significant mortality. Highly lethal systemic or oral infections have been useful in unravelling broad antiviral immune mechanisms (Dostert *et al.*, 2005; Wang *et al.*, 2006; Kemp & Imler, 2009; Nayak *et al.*, 2013; Karlikow *et al.*, 2014), but it is unlikely that the morbidity and mortality they cause is an accurate reflection of the level of disease experienced by flies in the wild, where viral infections appear to be widespread among many species of *Drosophila* as low level persistent infections with apparently little pathology (Kapun *et al.*, 2010; Webster *et al.*, 2015). Our understanding of the fitness costs of viral infection in *Drosophila* is therefore severely limited, which is striking given the evidence from population genetic data that viruses are major drivers of

adaptive evolution in *Drosophila* immune genes (Obbard *et al.*, 2006, 2009; Early *et al.*, 2016).

To gain a better understanding of the potential fitness costs of DCV infection, we measured the physiological and behavioural responses of flies challenged with DCV. We carried out two separate experiments, either challenging flies with a range of sub-lethal viral concentrations systemically through intra-thoracic injury (experiment 1) or exposing flies through the oral route of infection to a low, sub-lethal concentration of DCV (experiment 2). Our aim was not to compare the two routes of infection, but to address sub-lethal infections using both infection routes, as these are commonly employed in experimental infections. We focused on traits that have been previously shown to be affected by DCV infection such as survival, fecal excretion, and locomotor activity, as well as female reproductive output, which is ultimately important for evolutionary fitness. We find that even apparently benign, sub-lethal infections can cause significant physiological and behavioural morbidity that is relevant to fly fitness, and that these effects vary according to viral concentration, host genetic background and sex.

MATERIAL AND METHODS

Fly lines and rearing conditions

In experiment 1 (systemic DCV infection) we used *Drosophila melanogaster* line *G9a^{+/+}* described previously (Merkling *et al.*, 2015), kindly provided by R. van Rij (Radboud University, Nijmegen, NL). This line was maintained on standard Lewis Cornmeal medium (Lewis, 2014) under standard laboratory conditions at 25°C, 12h: 12h Light:Dark cycle. Experimental flies were generated by setting up 20 replicate Lewis vials with 15 males and 15 females to mate and lay eggs for 24 hours. Three-to-four-day-old adults that eclosed from the eggs laid during this period were infected systemically (see below) and then followed individually for health measures.

In experiment 2 (oral DCV exposure) we used ten *D. melanogaster* lines from the Drosophila Genetic Reference Panel (DGRP): RAL-83, RAL-91, RAL-158, RAL-237, RAL-287, RAL-317, RAL-358, RAL-491, RAL-732, and RAL-821. Given we had no prior knowledge of how the DGRP panel vary in response to oral DCV infection, these lines were chosen randomly. All lines were previously cleared of *Wolbachia* and have been maintained *Wolbachia*-free for at least 3 years. Fly stocks were kept at a density of 30 individuals in bottles on standard Lewis medium at 24.5± 0.5°C. Flies were allowed to mate and lay eggs for three days and then removed. When eggs had developed into three-day old imagoes, we picked 16 male and 16 female flies at random from each DGRP line (320 flies in total). Half of these flies (n=8 replicates) were individually exposed to DCV through the oral route of infection (see details below) and the other half were exposed to a sterile Ringers solution (7.2 g/L NaCl; 0.17 g/L CaCl₂; 0.37 g/L KCl,

diluted in sterile water, pH 7.4) as a control (n=8 replicates). Following infection, all flies were kept individually in vials kept in incubators at 24.5°C ± 0.5 with a 12h:12h light:dark cycle for the remainder of the experiment. Vials were randomized within trays to reduce any positional effects within incubators.

DCV stock and culturing

The Drosophila C Virus (DCV) isolate used in both experiments was originally isolated in Charolles, France (Jousset *et al.*, 1977), and was produced in Drosophila line 2 (DL2) cells as described previously (Longdon *et al.*, 2013; Vale & Jardine, 2015). Infectivity of the virus was calculated by measuring cytopathic effects in DL2 cells using the Reed-Muench end-point method to calculate the Tissue Culture Infective Dose 50 (TCID₅₀) (Reed & Muench, 1938). The DCV stock used in this experiment had an infectivity of approximately 4x10⁹ DCV infectious units (IU)/mL. This stock culture was serially diluted to achieve the desired concentrations (approximately 10² 10³ and 10⁵ DCV IU/mL for systemic infection and 10⁵ DCV IU/mL for oral infection) and kept at -80°C until needed.

Systemic DCV infection and viral titers

We exposed 20 individual male and female flies to each of 4 viral concentrations (160 flies in total)– 0 (control), 10², 10³ and 10⁵ DCV IU/ml, obtained by serial diluting the viral stock with 10mM Tris-HCl (pH 7.3). Flies were infected systemically by intra-thoracic pricking with a needle immersed in DCV suspension under light CO₂ anesthesia. Control flies were pricked with a needle dipped in sterile 10mM Tris-HCl (pH 7.3). An additional five individuals for each sex/dose combination were infected as described above to quantify DCV within

flies following infection, using the expression of DCV RNA. Flies were individually placed in TRI reagent (Ambion) following five days of infection (5 DPI), homogenized total RNA was extracted using Direct-zol RNA miniprep kit, which includes a DNase step (Zymo Research), reverse-transcribed with M-MLV reverse transcriptase (Promega) and random hexamer primers, and then diluted 1:2 with nuclease-free water. qRT-PCR was performed on an Applied Biosystems StepOnePlus system using Fast SYBR Green Master Mix (Applied Biosystems) and DCV primers, which include 5'-AT rich flaps to improve RT-PCR fluorescent signal (Afonina *et al.*, 2007) (DCV_Forward: 5' AATAAATCATAAGCCACTGTGATTGATACAACAGAC 3'; DCV_Reverse: AATAAATCATAAGAAGCAGATACTTCTTCCAAACC). We measured the relative fold change in DCV RNA relative to *rp49*, (Dmel_rp49 Forward: 5' ATGCTAAGCTGTCGCACAAATG 3' ; Dmel_rp49 Reverse: 5' GTTCGATCCGTAACCGATGT 3'). an internal *Drosophila* control gene, calculated as $2^{-\Delta\Delta Ct}$ (Livak & Schmittgen, 2001).

Oral DCV exposure

In separate pilot infections, we determined that a DCV culture diluted to contain approximately 10^5 DCV RNA copies was enough to establish a viable infection (Figure S1), but did not cause noticeable mortality, and we used this dilution of DCV stock to inoculate all ten DGRP lines. Individual flies were exposed to DCV in vials containing Agar (5% sugar) using 3mL plastic atomizer spray bottles containing 2mL of the sub-lethal DCV dilution. One spray, releasing roughly 50 μ L of DCV dilution (or sterile Ringer's solution), was deployed into each vial. Flies

were left in the these 'exposure vials' for three days to allow them to ingest the viral solution during feeding and grooming, and then tipped into vials containing clean, blue-dyed Lewis medium (see below).

Survival following infection

Both systemically and orally infected flies were housed individually following infection in vials containing Lewis medium. In the systemic infection experiment, flies were monitored daily for mortality for 38 days post-infection and were transferred to fresh food vials once a week. In the oral infection experiment, flies were transferred to fresh food vials every 3-4 days, and mortality was recorded at this point for the first 32 days post infection and then daily until 40 DPI (oral infection).

Fecal excretion following oral DCV exposure

Following the exposure period, flies were tipped into vials containing blue-dyed Lewis medium. Blue medium was prepared by adding 0.5g/L FIORI COLORI brilliant blue FCF E133 granules to standard Lewis medium. Flies remained on blue Lewis food for the remainder of the experiment and were tipped to new blue Lewis vials every three to four days. When flies were tipped to new vials, the old vials were kept for fecal spot counts (measured immediately) and fecundity measures (see below). Fecal spots were recorded by photographing vials with a Leica S8APO microscope. A slip of white printer paper (2.5cm x 8.5cm) was inserted into each vial to ensure only spots on one side of the vial were being photographed. These images were then analyzed with ICY image software (Version 1.6.1.1 ICY - Bio Imaging Analysis) and fecal spots were

counted using 'spot detection' analysis on a 2cm x 4cm region of interest. Each image was checked individually for miscounts, and miscounted spots were removed. Fecal excretion was recorded for 30 days following infection.

Fecundity

All fecundity estimates are based upon mating that occurred before infection during the first 3-4 days after eclosion. The fecundity of individual flies was measured by counting viable offspring emerging in the vials they were reared in, which happened weekly until day 30 post infection in the systemically infected flies, and every 3-4 days in the orally infected flies, for 28 days following exposure to DCV. Short-term fecundity estimates have been shown to be well correlated with lifetime reproduction in *D. melanogaster* (Nguyen & Moehring, 2015). Vials that individuals were tipped from (and following the recording of fecal shedding in the oral infection experiment), were placed in the incubators at 24.5°C ± 0.5 with a 12h:12h light:dark cycle to allow any offspring to develop. After 14 days, the total number of living emerged adult offspring within each vial was recorded as a measure of female fecundity.

Activity

Locomotor activity was measured using the Drosophila Activity Monitor (DAM2, Trikinetics) as described previously (Pfeifferberger *et al.*, 2010; Vale & Jardine, 2015). In the DAM, individual fly activity is recorded when individually housed flies break an infrared beam passing through a transparent plastic tube placed symmetrically inside a DAM unit. In systemically infected flies, as we used females to measure fecundity (see above), activity was measured on 10 replicate

male flies for each DCV dose (40 flies in total), starting the day following septic injury, and measured for 2 weeks following infection. In the oral infection experiment, activity was recorded for 24 hours, fourteen days after the initial oral exposure. These differences in the timing of activity measurements arise from the faster and more severe effects of systemic infections on locomotor behavior, while we have found that effects on activity following oral infection take longer to manifest, and become apparent 10-15 days after DCV ingestion(Vale & Jardine, 2015). Four replicate flies for each DGRP (10 lines) / sex (M/F) / infection (DCV/Control) combination were tested (160 flies in total). In both experiments, flies were placed individually in a single DAM tube containing a small agar plug on one end, and allocated a slot in one of five DAM unit (each unit can house a maximum of 32 tubes). At least one slot in each DAM unit was filled with an empty tube and at least two slots were left empty as negative controls. All DAM units were placed in the incubator (25 °C 12:12 light:dark cycle) and continuous activity data was collected every minute for 24 hours. Raw activity data was processed using the DAM System File Scan Software (www.trikinetics.com) and the resulting data was manipulated using R v. 3.1.3 (The R Foundation for Statistical Computing, Vienna, Austria). Flies that died during the DAM assay (6/40 flies in the systemic infection experiment; 25/160 in the oral infection experiment) were removed from the analysis because they would wrongly bias the estimate of activity.

Data analysis

All analyses were carried out in JMP 12 (SAS). Survival data was analyzed on the 'day of death' using a Cox Proportional Hazards models in with 'fly sex' and 'DCV

exposure' and their interaction as fixed effects (systemic infection experiment) or 'fly sex', fly 'line' and 'DCV dose' and their interactions as fixed effects (oral infection experiment). In the systemic infection, DCV titers were Log₁₀-transformed and analyzed in a linear model with 'DCV Dose' and 'Sex' and their interaction as fixed effects. Fecundity following systemic infection was calculated on the cumulative number of emerged offspring in a model containing 'DCV dose' as a fixed effect. In the oral exposure experiment, the cumulative number of offspring was analyzed in a model including 'Fly line' and 'DCV exposure' and their interaction as fixed effects. Total excretion per fly was analyzed using a linear model with 'Fly line', 'DCV exposure', and 'sex' as categorical fixed effects, 'Time' as a continuous covariate, and all pair-wise interactions. Activity was analyzed as the total number of DAM beam breaks recorded per day. Activity following systemic infection was analyzed in a linear model with 'DCV dose' and 'Time' as fixed effects. Activity following oral infection was measured for 24h and analyzed in a linear model with 'Fly line', 'Sex' and 'DCV exposure' as fixed effects. In all analyses, individual replicate was included as a random factor, and in all cases accounted for only 2-5% of the total variance.

RESULTS

Experiment 1: Sub-lethal systemic infection

In a first experiment, we tested how systemic infection with very low concentrations of DCV (10^2 , 10^3 and 10^5 DCV IU/ ml) affected fly health. We have previously observed that DCV is able to establish and grow when inoculated into flies at these low doses (Figure S2). The survival of both female and male flies exposed to doses of 10^2 and 10^3 DCV IU/ ml did not differ from control flies that had been pricked with sterile buffer solution (Figure 1a). In females, 100% flies exposed to these doses survived infection during the 38-day survival assay, while roughly 20% of males died during this period (Figure 1a). However, this difference in survival between sexes ('sex' effect, Table 1), was also observed in control flies and therefore is likely to reflect sex-specific responses to injury during intra-thoracic pricking than to infection. Flies infected with a slightly higher concentration of 10^5 DCV IU/ ml died significantly faster than control flies. This virus concentration-specific pattern of mortality was generally consistent with the observed DCV titers measured 5 days following infection, (Table 2, 'dose' effect) which were generally higher in male flies across all DCV concentrations (Table 2, 'sex' effect, Figure 1b). Our experiment therefore spanned the range of sub-lethal viral doses, with 10^5 DCV IU/ ml being the lowest virus concentration with lethality in the experiment (Figure 1a).

Fecundity following systemic DCV infection

We used mated females, which allowed us to quantify fly reproductive health during systemic infection by following the number of adult offspring produced by individual females for 30 days following infection. The total fecundity

measured during this period varied according to the dose females had received ($F_{3,66} = 10.32$, $p < 0.0001$) and we observed that the total reproduction of infected flies was higher than control flies, and increased in a dose-specific manner (Figure 1c).

Activity following systemic DCV infection

The locomotor activity of individual male flies infected systemically with all sub-lethal concentrations of DCV was measured during 18 days after infection in a Trikinetics® Drosophila Activity Monitor (DAM). All flies included in the analysis remained alive for the whole period, so changes in activity were not confounded with potential death of individual flies. We found that flies in all treatments, including uninfected controls, showed a reduction in activity over the course of the activity assay (Figure 1d, Table time effect). This general effect is not especially surprising given the constrained environment experienced by flies in the DAM tubes, and that the only source of nutrition and hydration is small agar plug. However, our analysis showed that the temporal reduction in activity depended on the dose that flies had received ('time x dose' interaction, Table 1). In the early stages of infection flies receiving the higher of the 4 doses (10^3 and 10^5 DCV copies) showed a reduction in activity relative to control flies and those receiving the lowest dose. Over time, a reduction in locomotor activity was most apparent in flies infected with the highest dose of 10^5 DCV copies (Figure 1d).

Experiment 2: Sub-lethal gut infection

In a separate experiment, we tested how exposure to a single sub-lethal dose of DCV through the oral route of infection impacted upon fly health. We conducted

the experiment on ten fly lines from the DGRP panel (Mackay *et al.*, 2012) and we included both male and female flies to test for the effects of host genetic background and sex in response to sub-lethal oral infection. While DGRP lines differ in their lifespan in the absence of infection (Durham *et al.*, 2014), we did not detect any difference between DGRP lines or between sexes in their survival during oral DCV infection compared to control flies (Table S1) which, as expected, was generally non-lethal across all lines.

Fecundity following oral exposure to DCV

Despite not observing any effects on fly survival during infection, we detected significant variation in reproductive health following exposure to DCV. The total fecundity of females during the 28 days following oral exposure to DCV (or a control inoculum) varied significantly between DGRP lines (Figure 2; Table 2), reflecting well-known genetic differences in the lifetime reproductive output of these lines (Durham *et al.*, 2014). In addition, we found line-specific fecundity responses to DCV infection ('infection status x line', Table 2, see also Table S2 for pairwise contrasts). In some lines (158, 491, 317) low-level oral infection resulted in a decrease in fecundity; in other lines (821, 358) there was no detectable effect of DCV exposure; while in 2 lines we detected significant increases in fecundity in DCV infected flies compared to uninfected control flies of the same genetic background (Figure 2; see Table S2 for least-square pairwise contrasts).

Locomotor activity following oral exposure to DCV

Overall, DGRP lines differed in their activity in a sex specific way ('Fly line x Sex' effect Table 2), but these differences were not altered by infection. While we detected a reduction in locomotor activity following systemic infection (Figure 1d), we did not detect any effect of oral DCV exposure on the overall activity of flies (Table 2, Figure 3).

Fecal excretion following oral exposure to DCV

We quantified fecal excretion for 30 days following DCV exposure as a proxy for gut health, by counting fecal spots excreted into vials after ingestion of blue-dyed food. Overall we found that males showed higher levels of fecal excretion compared to females (Table 2, 'sex' effect; Figure 4) and that DCV infection was associated with a general reduction in fecal excretion throughout the 30-day observation period ('Infection status' effect, Figure 4). However, we found that males and females differed in the overall severity of this reduction ('sex x infection status' effect), with males showing a greater reduction in defecation overall (Figure 4). Furthermore, we found significant variation among the DGRP lines in the magnitude of the effect of DCV on fecal excretion ('fly line x infection status' effect).

DISCUSSION

We find that sub-lethal infections with DCV can cause measurable morbidity that is relevant for the fitness costs experienced by *D. melanogaster* during DCV infection. In two independent experiments using sub-lethal concentrations of either systemic or oral DCV infections, we observed effects on fly reproductive output, digestive health, and locomotor activity.

Systemically infected flies increase reproductive output

We found that the fly line used in the systemic infection experiment showed an increase in reproductive output when infected with sub-lethal doses of DCV. There are numerous examples from both invertebrates and vertebrates of fecundity increases following infection (Bonneaud *et al.*, 2004; Vale & Little, 2012; Leventhal *et al.*, 2014; Vézilier *et al.*, 2015). In addition, earlier work reported that DCV infection could increase ovariole number and decrease development time in *D. melanogaster* (Thomas-Orillard, 1984; Gomariz-Zilber & Thomas-Orillard, 1993). However, a subsequent re-analysis of these data showed very weak support for the beneficial effects of DCV infection (Longdon, 2015). It is notable however that neither of the earlier studies measured the number of viable offspring of infected flies compared to healthy ones. The fecundity data we report therefore suggests that DCV may indeed result in increased reproductive output.

A dose-dependent increase in fecundity could suggest a direct effect of DCV infecting fly ovaries, but it is unclear why such a strategy would be adaptive for

the virus. An alternative hypothesis may instead involve more complex interactions between the allocation of resources during DCV infection, and how they relate to fly nutritional stress and reproductive investment. For example, *D. melanogaster* females selected under conditions of nutritional stress were found to produce a greater number of ovarioles, while the offspring of starved mothers also exhibited greater investment in reproduction (Wayne *et al.*, 2006). Similar to the studies described above (Thomas-Orillard, 1984; Gomariz-Zilber & Thomas-Orillard, 1993), this work also focused on ovariole number and egg production, and did not quantify female lifetime fecundity. Given that DCV infection is known to lead to intestinal obstruction, one possibility for the increase in the number of adult offspring we observed in infected flies is that DCV-induced nutritional stress leads to a greater production of ovarioles, and consequently, an increased number of offspring. Given we only tested a single fly line however, it is important to note that this response may not be universal. As we discuss below fecundity responses to infection have generally been found to differ between host genotypes (Vale & Little, 2012; Parker *et al.*, 2014)

Fecundity costs and benefits of DCV infection are genotype-specific

Similar to systemically infected flies (Figure 1c), we also find evidence for fecundity benefits in orally exposed flies, but these benefits were only revealed in two out of the ten genetic backgrounds we tested. Indeed, in three of the tested lines, DCV infection resulted in lower reproductive output. Taking fecundity as a proxy for evolutionary fitness, the existence of genotype specific fitness costs and benefits means that DCV could be a potentially powerful driver of *D. melanogaster* evolutionary dynamics. Previous analyses of *Drosophila* spp.

population genetic data have shown that the fastest evolving *D. melanogaster* genes are those involved in RNAi-based antiviral defense (Obbard *et al.*, 2006, 2009; Early *et al.*, 2016), but the DCV-induced fitness costs that drive this rapid evolution in wild-infected flies (where infections are persistent and often non-lethal), has remained obscure. These data suggest that genotype-specific fecundity costs and benefits of DCV infection could potentially mediate the arms-race between flies and viruses.

Systemically infected flies show a dose-dependent decline in activity over time

Reduced activity, or lethargy, following infection is a common response to infection across a range of taxa (Hart, 1988; Adelman & Martin, 2009; Sullivan *et al.*, 2016). The most obvious explanation for reduced activity is simply that infected individuals are sick, and lethargy reflects the underlying pathology of infection (Moore, 2013). A popular alternative explanation is that infection-induced lethargy evolved as an adaptive host strategy that conserves energy, which may then be allocated to other physiological tasks such as mounting an immune response (Hart, 1988; Adelman & Martin, 2009).

Support for the adaptive nature of these ‘sickness behaviours’ has come mainly from vertebrate species challenged with deactivated pathogens or their derived components, which are sufficient to stimulate an immune response without causing pathology (Adelman & Martin, 2009; Lopes *et al.*, 2016). In addition to vertebrates, sickness behaviors including lethargy and anorexia have also been described in insect hosts (Ayres & Schneider, 2009; Kazlauskas *et al.*, 2016; Sullivan *et al.*, 2016). However, in the current experiment it is not possible to

disentangle the effect of an adaptive sickness behavior from the direct effect of pathology caused by replicating DCV. Regardless of the underlying cause of reduced activity, it is likely to come at an additional cost of lower involvement in fitness-enhancing activities such as foraging, competing for resources with conspecifics, or courtship and mating (Adelman & Martin, 2009; Adamo *et al.*, 2015; Vale & Jardine, 2016). Further, reduced activity following infection can also reduce the potential for disease spread (Lopes *et al.*, 2016). In the context of understanding sub-lethal DCV infection in an ecological setting, reduced activity may therefore be a potentially important source of DCV-induced fitness costs and benefits.

We did not find an effect of oral DCV exposure on fly activity. Previous work has shown that *Drosophila*, especially females, show a reduction in activity following oral infection with DCV (Vale & Jardine, 2015). However, the viral concentration that flies were exposed to in that experiment was at least 1000x higher, so it is likely that in the current experiment flies did not ingest virus in quantities large enough to affect locomotor activity.

The severity of DCV-induced digestive dysfunction is sex-specific

Previous work has shown that DCV infection results in digestive dysfunction, leading to increased body mass due to the inability to excrete digested food (Arnold *et al.*, 2013; Chtarbanova *et al.*, 2014). We found that this measure of gut health varied between genotypes and also between sexes. Extensive genetic variation for gut immune-competence has previously been reported in the DGRP panel (Bou Sleiman *et al.*, 2015), which could underlie some of the variation we

observe in DCV-associated digestive dysfunction in some lines. Although that study focused on enteric infection with entomopathogenic bacteria, the mechanisms that mediate variation in gut health during infection include general processes of gut damage and repair, such as the production of reactive oxygen species (ROS) and the production of intestinal stem cells during epithelial repair (Buchon *et al.*, 2013). It is plausible that these mechanisms also mediate disease severity during enteric virus infection, but we are unaware of any systematic study of genetic variation in gut immune-competence during viral infection.

The mechanistic basis of the observed sex differences in fecal excretion is less clear. The Malpighian tubules are the main organ involved in osmoregulation and excretion of waste matter in insects (Dow & Davies, 2001). *D. melanogaster* male and female Malpighian tubules have been shown to differ at the transcriptional level with over 18% of genes (2308 genes) showing sex-specific expression (Huylmans & Parsch, 2014). We measured fecal excretion by quantifying fecal spots on the sides of the vials. Given that females are known to also spend more time feeding (Wong *et al.*, 2009), it is possible that females also defecate more on the surface of the food compared to males, and therefore spend less time on the sides of the vials. Only a few studies have investigated sex differences in fecal excretion in *D. melanogaster*, finding inconsistent patterns of excretion between sexes (Zeng *et al.*, 2011; Urquhart-Cronish & Sokolowski, 2014). The link between fecal excretion and fitness is not as clear as with fecundity or locomotor activity, but it is relevant in the context of disease transmission of fecal-orally transmitted pathogens such as DCV. The study of temporal trends in fecal excretion and how they vary with host sex and genetic background may

therefore be used as a useful model to understand the sources of heterogeneity in pathogen shedding (Vale *et al.*, 2013).

Concluding remarks

Altogether, these measures of sub-lethal morbidity give insight into the potential fitness costs of low-level, persistent DCV infection in *Drosophila*. More generally, the combination of both positive and negative effects on fly fitness effects according to the specific host genetic background presents a non-trivial evolutionary scenario for host immune defense (Gandon & Vale, 2014). For instance, frequent encounters between beneficial symbionts and detrimental pathogens are hypothesized to have played a role in the evolution of aphid immune systems, which lack several components of the IMD immune pathway critical for the recognition and elimination of Gram-negative bacteria (Gerardo *et al.*, 2010). The combination of fitness costs and benefits of infection, such as those incurred during DCV infection, may therefore have driven the evolution of immune defense across a wide range of host taxa, from insects to mammals (Elsik, 2010; Gerardo *et al.*, 2010; Lee & Mazmanian, 2010; Gandon & Vale, 2014).

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528

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Figure legends

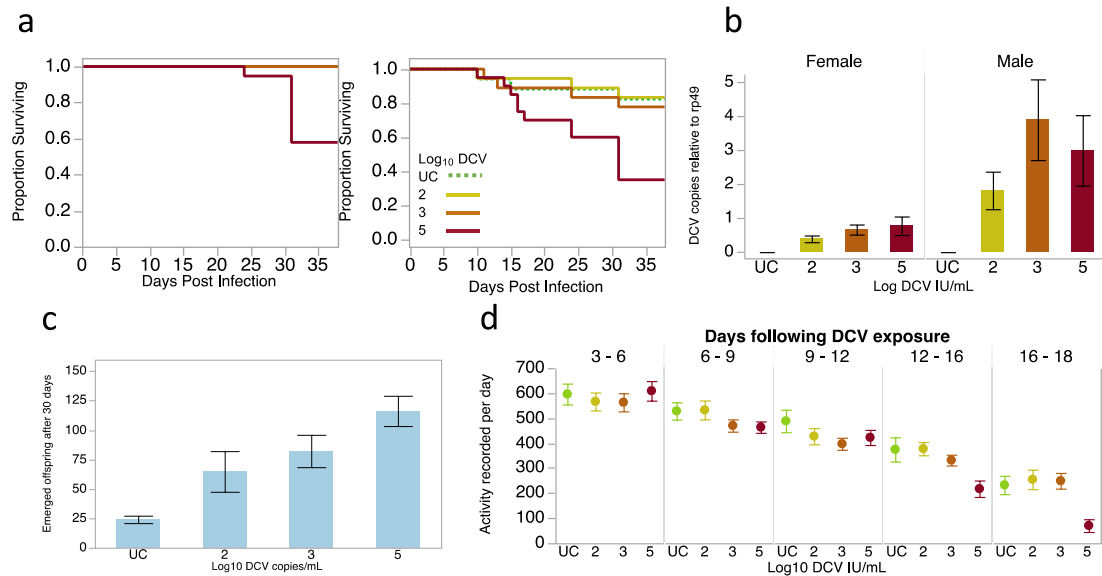
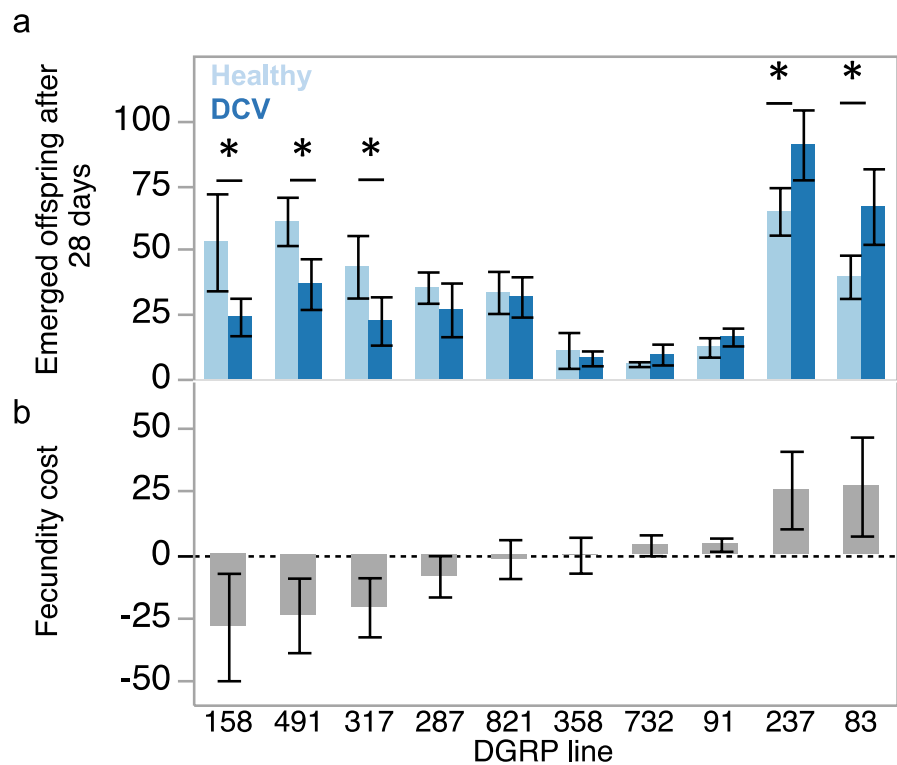


Figure 1. Sub-lethal systemic infection. 1a. Kaplan-Meier curves showing the survival of 20 replicate flies exposed systemically to sub-lethal concentrations of DCV. 1b. DCV titers measured in male and female flies relative to an internal control gene (*rp49*), following 3 days of systemic infection with sub-lethal concentrations of DCV. For each DCV concentration, data are the average of duplicate qPCR reactions for 5 individual flies. 1c. The total number of emerged adult offspring recorded for 30 days following systemic infection based on mating that occurred before infection during the first 3-4 days after eclosion. Data are the means \pm SE of 18-19 replicate female flies. 1d. Daily locomotor activity of male flies following systemic infection with DCV. Data are 3 day averages of 7-10 replicate flies for each inoculation concentration. UC are uninfected controls.

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735 **Figure 2.** Fecundity following oral DCV exposure. 2a. The cumulative number of
736 adult offspring from healthy (light bars) or DCV-exposed (dark bars) single
737 female flies over the course of the 28-day experiment. 2b. Shows the fecundity
738 difference between healthy and infected flies for the same 10 DRGP lines. In both
739 plots, DRGP lines are ordered from the greatest decrease to the highest fecundity
740 increase. Significant pairwise contrasts (reported in Table S2) are indicated by
741 asterisks. Data are the mean \pm SE of eight individual replicate females.

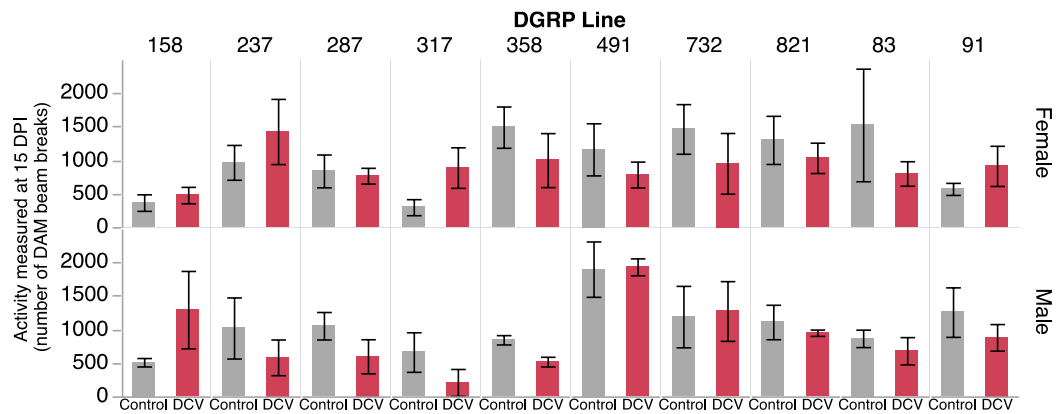


Figure 3. Locomotor activity following oral DCV exposure. Data show mean \pm SE activity of four replicate flies per sex and DGRP line, measured for 24 hours 14 days following exposure to DCV (red) or uninfected controls (grey).

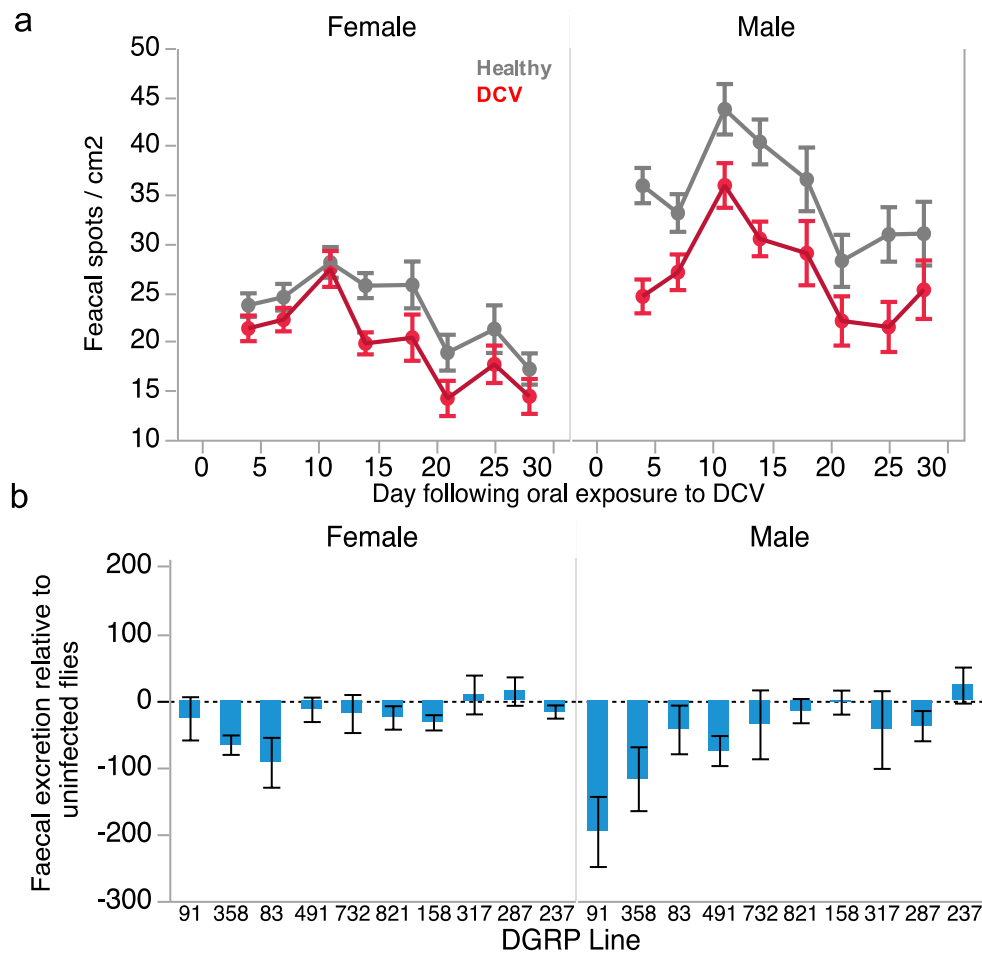


Figure 4. Fecal excretion following oral DCV exposure. 4a. The general effect of DCV exposure (red) or a control inoculum (grey) on the number of fecal spots shed over time. Data are plotted separately for males and females. Each time point is the mean \pm SE of 8 replicate individual flies averaged across all 10 DGRP lines. 4b. Shows the difference between control and infected flies for each DRGP line. Data are the mean \pm SE of eight individual replicate flies for each sex and line combination.

Table 1 - Systemic infection

<i>Survival</i>	DF	χ^2	p-value
DCV concentration	4	45.24	0.0001
Sex	1	8.37	0.0038
DCV concentration \times Sex	2	8.26	0.0161
<i>Viral titer</i>	DF	F Ratio	p-value
DCV concentration	3	3.14	0.0399
Sex	1	7.34	0.0111
DCV concentration \times Sex	3	1.35	0.2776
<i>Activity per day</i>			
Time (DPI)	1	290.68	0.0001
DCV concentration	3	5.17	0.0016
Time (DPI) \times DCV concentration	3	5.51	0.001

Table 2 - Oral infection

	DF	F Ratio	p-value
<i>Fecundity</i>			
DGRP Line	9	16.17	<.0001
DCV exposure	1	0.99	0.3186
DGRP Line × DCV exposure	9	2.59	0.0076
<i>Activity per day</i>			
DGRP Line	9	2.91	0.0037
Sex	1	0.02	0.8947
DCV exposure	1	1.45	0.2315
DGRP Line × Sex	9	2.18	0.0277
DGRP Line × DCV exposure	9	0.67	0.7352
Sex × DCV exposure	1	0.12	0.7244
<i>Fecal excretion</i>			
DGRP Line	9	32.17	0.0001
Sex	1	212.66	0.0001
Time (DPI)	1	29.95	0.0001
DCV exposure	1	72.83	0.0001
DGRP Line × DCV exposure	9	4.46	0.0001
Sex × DCV exposure	1	13.45	0.0003
Time (DPI) × DCV exposure	1	0.23	0.6295
DGRP Line × Sex	9	31.22	0.0001
DGRP Line × Time (DPI)	9	1.28	0.2405
Sex × Time (DPI)	1	0.06	0.806

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Supplementary File for

Costs and benefits of sub-lethal *Drosophila C* Virus infection

This file contains:

- Table S1. Cox proportional hazards analysis of survival following oral exposure to DCV.
- Table S2. Least Square Means Student's t pairwise contrasts between exposed and control fecundity following oral DCV exposure.
- Figure S1. DCV increases in titer following oral exposure to approximately 10^5 DCV copies.
- Figure S2. DCV increases in titer following systemic challenge with 10^2 , 10^3 and 10^5 DCV IU/ ml.

Table S1. Output of Cox proportional hazard model testing variation in survival following oral infection.

Survival during oral infection	DF	χ^2	p-value
DGRP Line	9	3.87084122	0.9197
Sex	1	3.82E-07	0.9995
DGRP Line*Sex	9	2.85864198	0.9696
Infection status	1	4.73E-08	0.9998
DGRP Line* Infection status	9	0.74383375	0.9998
Sex* Infection status	1	1.07E-06	0.9992
DGRP Line*Sex* Infection status	9	0.25051421	1

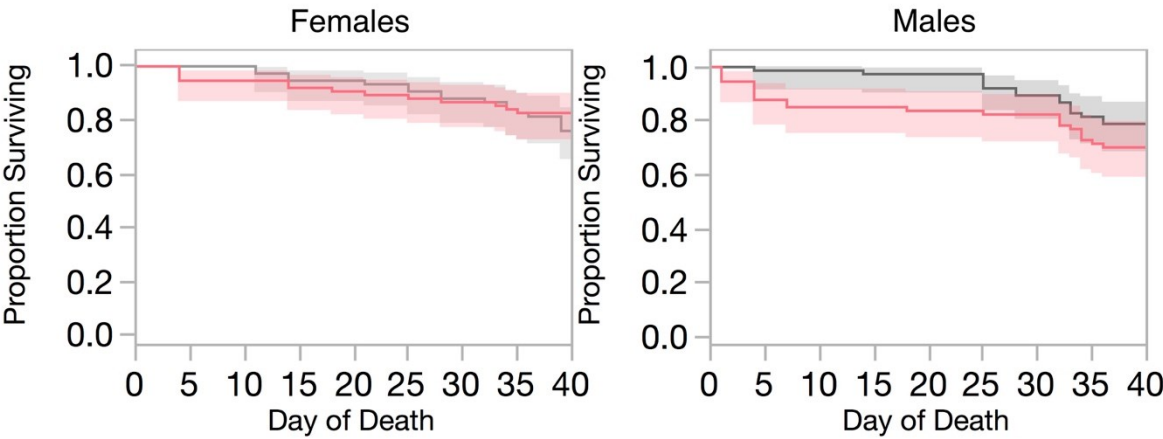


Table S2. Least Square Means Student's t pairwise contrasts between exposed and control fecundity following oral DCV exposure

DGRP line	NumDF	F Ratio	p-value
83	1	5.0178	0.036
91	1	0.2916	0.590
158	1	7.4368	0.007
237	1	5.6287	0.019
287	1	1.0515	0.306
317	1	4.6993	0.042
358	1	0.0525	0.819
491	1	4.7059	0.031
732	1	0.3253	0.569
821	1	0.0813	0.776

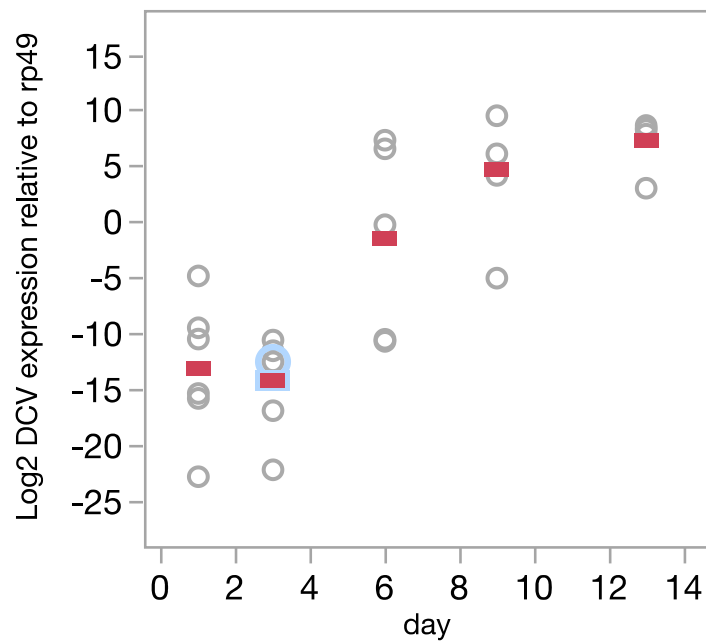


Figure S1. DCV increases in titer following oral exposure to with approximately 10^5 DCV copies ($F_{1,27} = 57.97$, $p < 0.001$). This experiment was carried out in *D. melanogaster* OreR. Data show the Log2 DCV expression relative to an internal *Drosophila* control gene (rp49), measured in six individual female flies at each time point following exposure. Oral exposure to DCV was carried out as described in the main text.

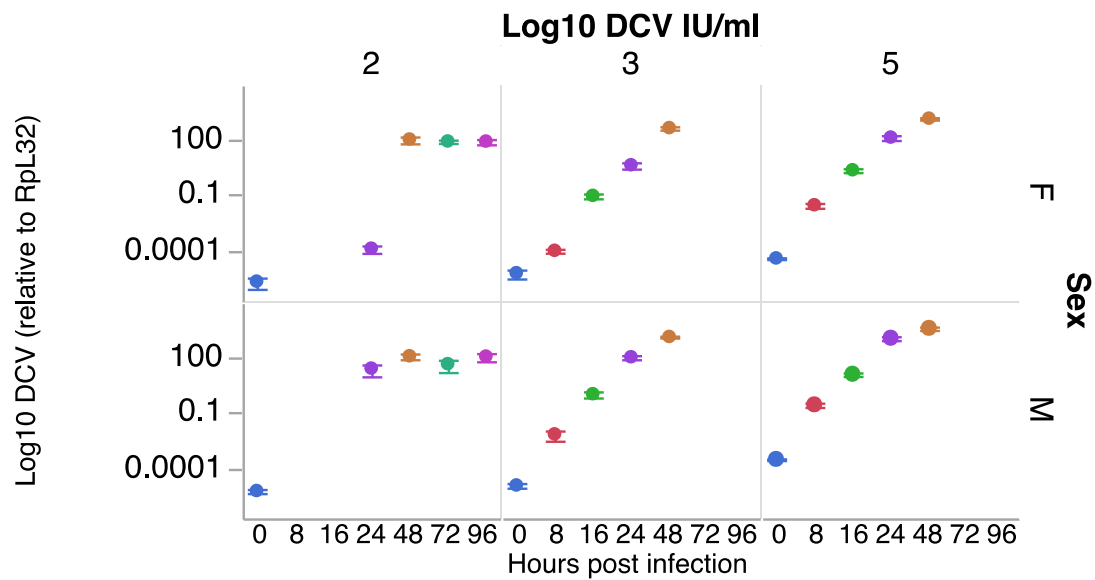


Figure S2. These data show DCV expression relative to the internal control gene Rpl32 measured at roughly 8-hour intervals. Male (M) of female (F) *D. melanogaster* (Oregon R, Wolbachia-negative) were challenged with 2, 3 or 5 Log10 DCV IU/ml. Data show means \pm SE of duplicate qPCRs on 3 replicate groups of 5 flies per sex/DCV concentration.